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(54) **ANCHORS FOR BODILY IMPLANTS AND METHODS FOR ANCHORING BODILY IMPLANTS INTO A PATIENT'S BODY**

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A61F 2/00 (2006.01)

(52) **U.S. Cl.**
CPC **A61F 2/0045** (2013.01); **A61F 2210/0004** (2013.01); **A61F 2220/0008** (2013.01)

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CPC ... A61F 2/0045; A61F 2/0063; A61F 2/0811; A61F 2220/0008; A61B 17/0401; A61B 2017/00805; A61B 2017/0409; A61B 17/0487
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See application file for complete search history.

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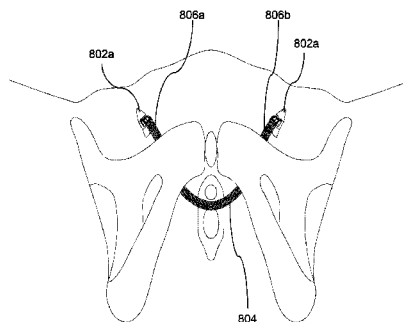
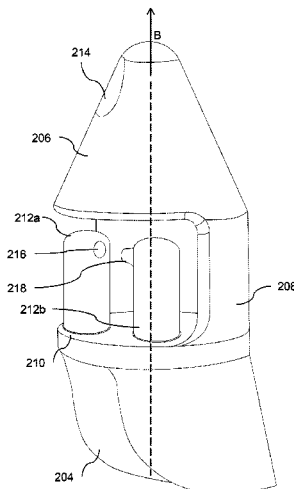
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(57) **ABSTRACT**

An anchor is provided for anchoring a bodily implant within a body of a patient. The anchor includes an implant engaging portion for engaging the bodily implant, wherein the implant engaging portion is disposed on a lateral portion of the anchor. The anchor further includes a distal end portion configured to pass through a passageway in the patient's body, the passageway defining a first axis and a proximal end portion disposed longitudinally opposite to the distal end portion on the anchor. The anchor defines a second axis extending from the distal end portion to the proximal end portion. The anchor is configured to rotate when a force is applied to the bodily implant such that the second axis defined by the anchor forms an angle with the first axis defined by the passageway.

19 Claims, 15 Drawing Sheets



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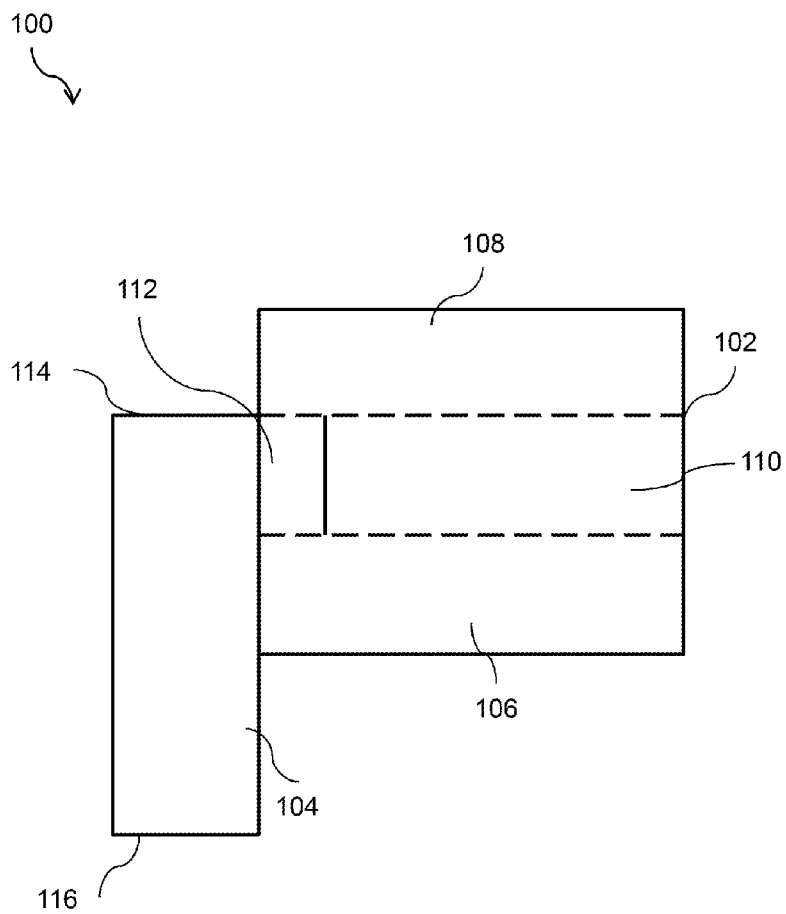


Fig. 1

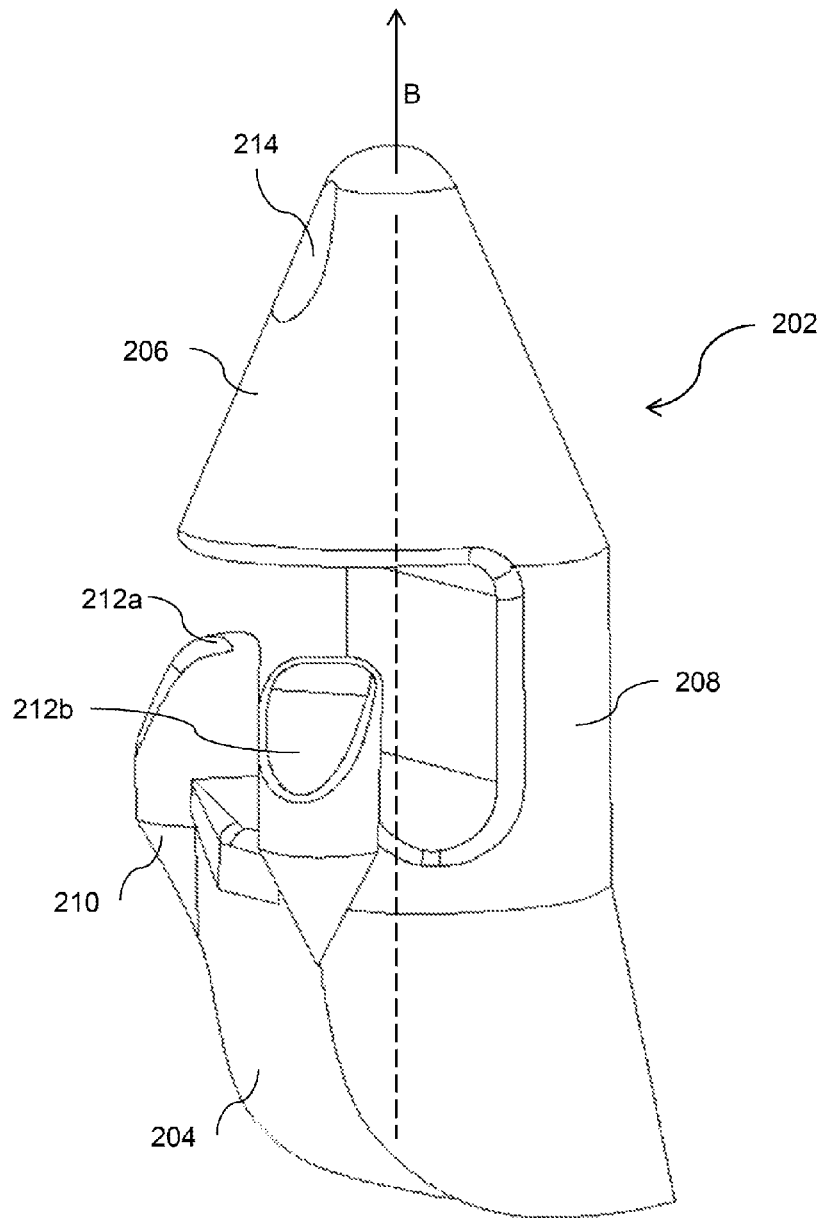


Fig. 2A

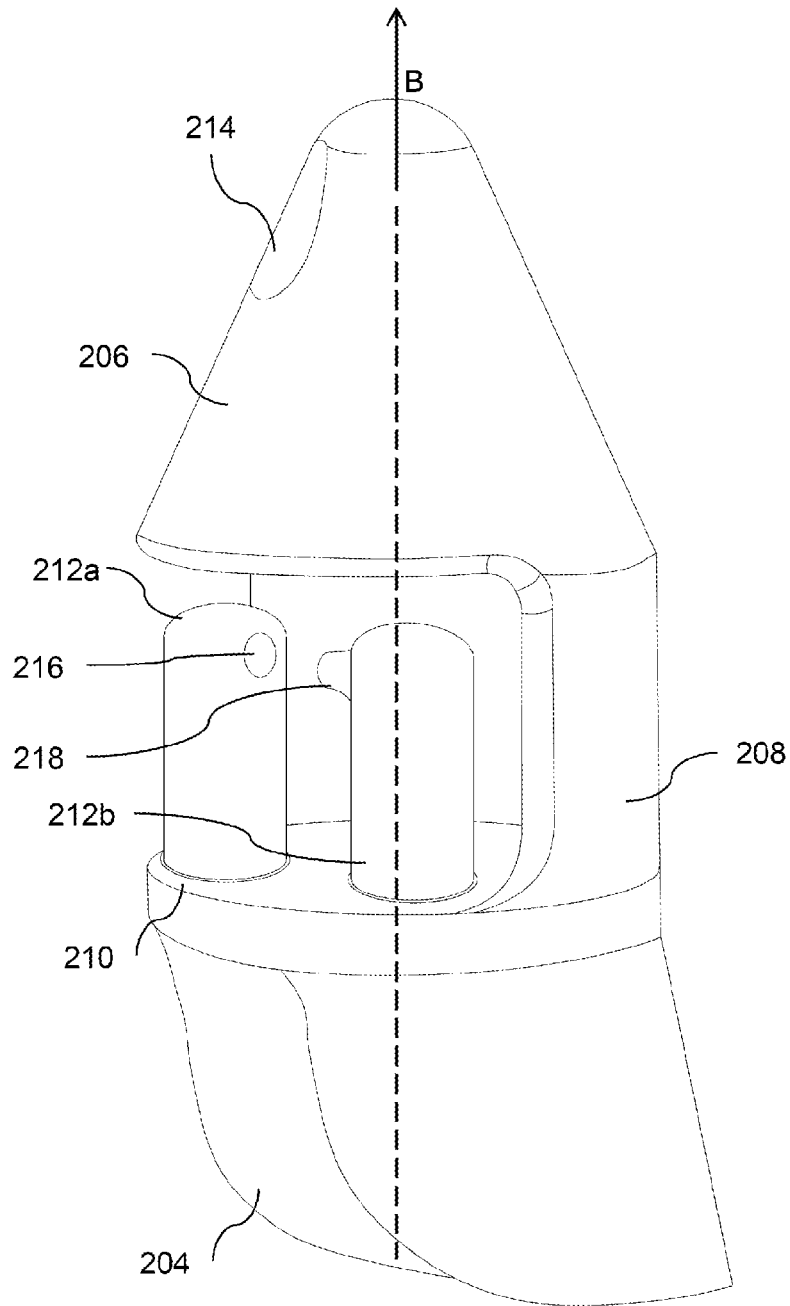


Fig. 2B

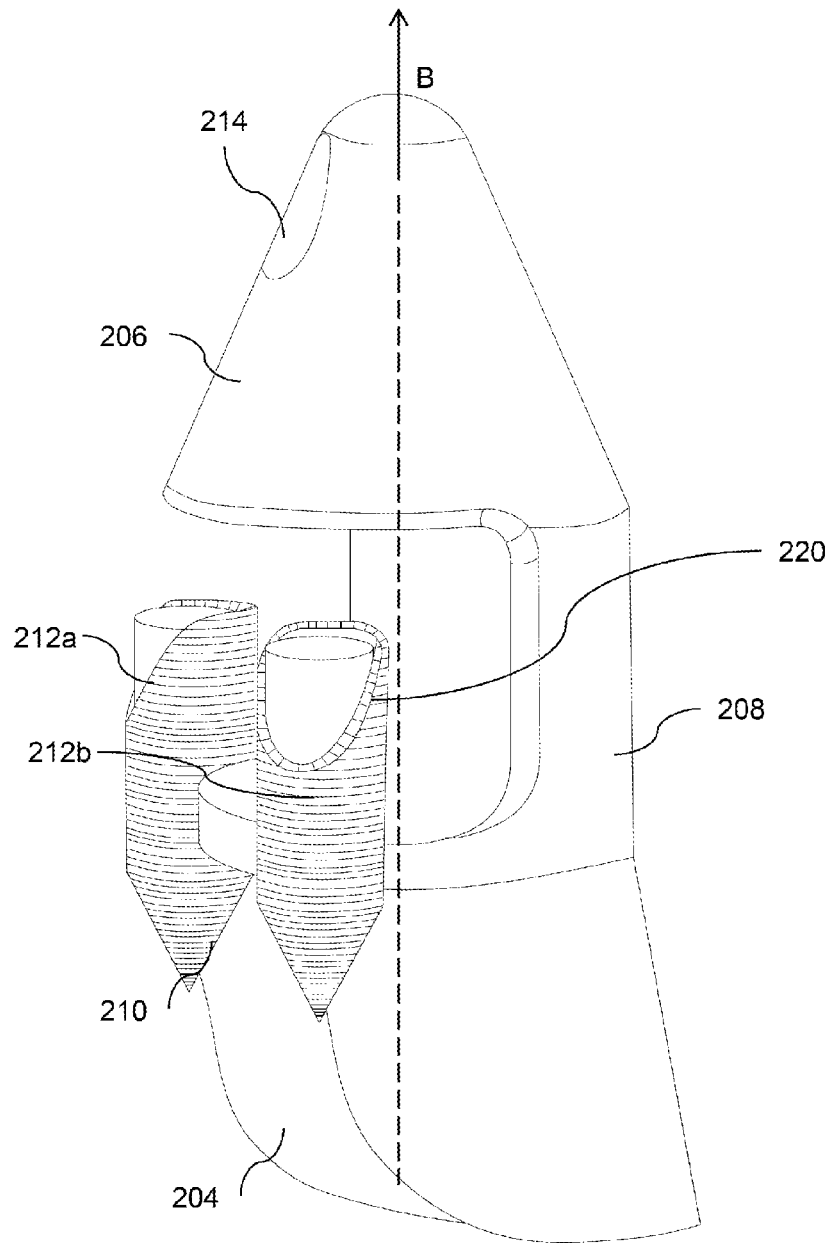


Fig. 2C

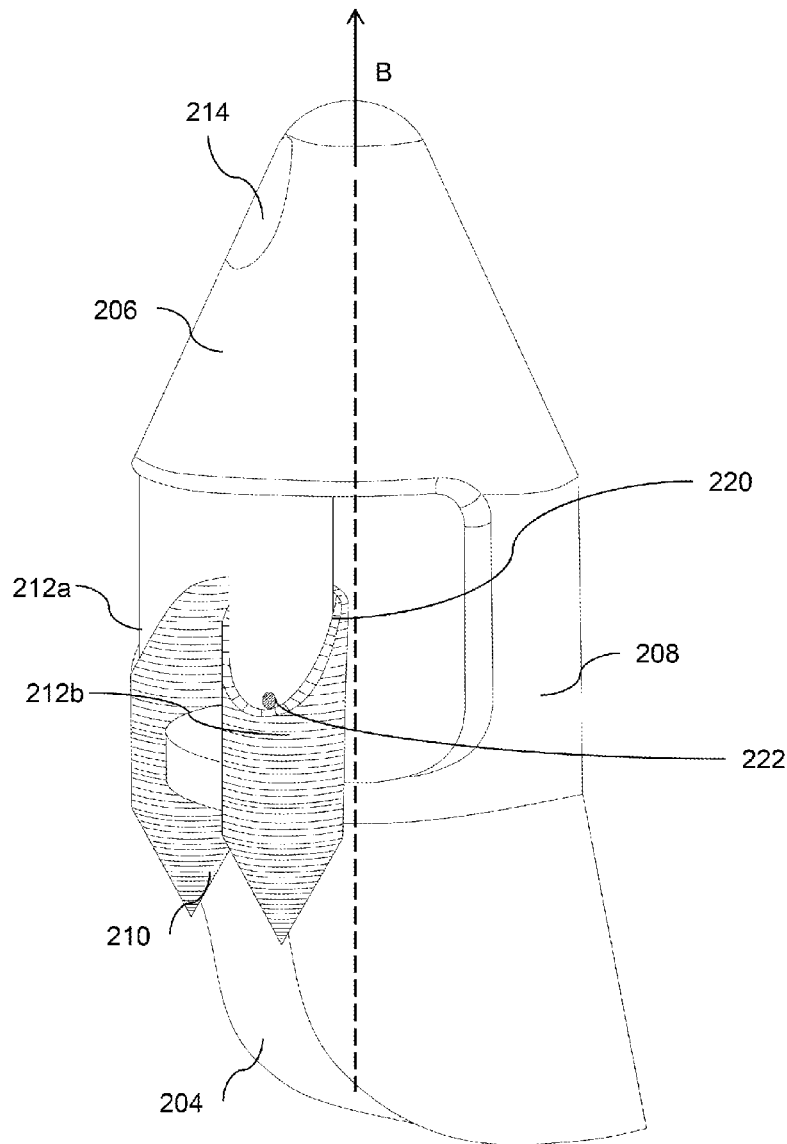


Fig. 2D

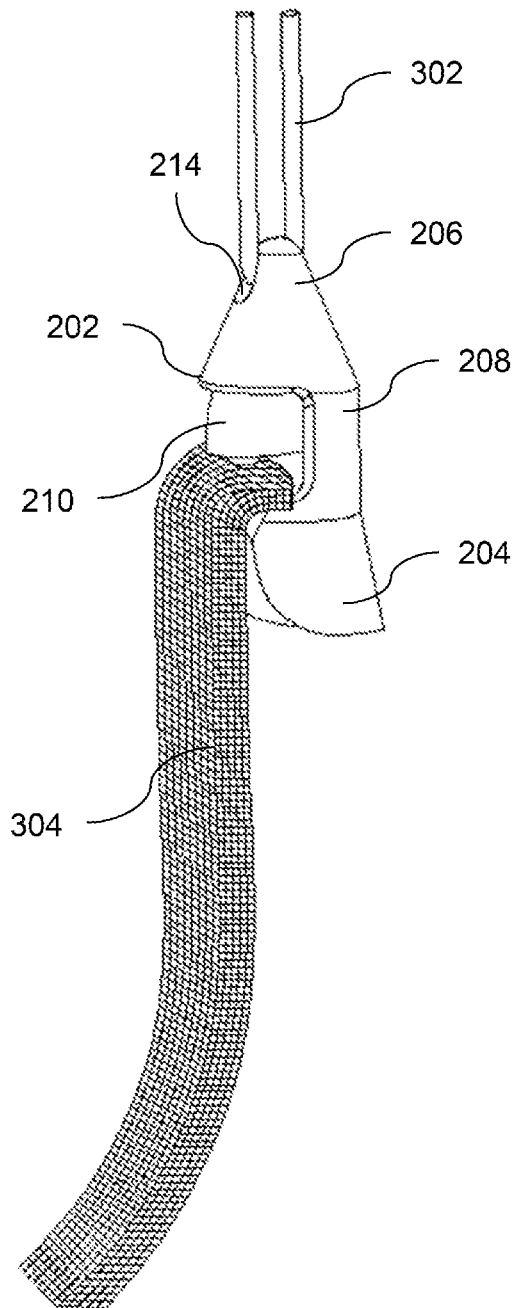


Fig. 3

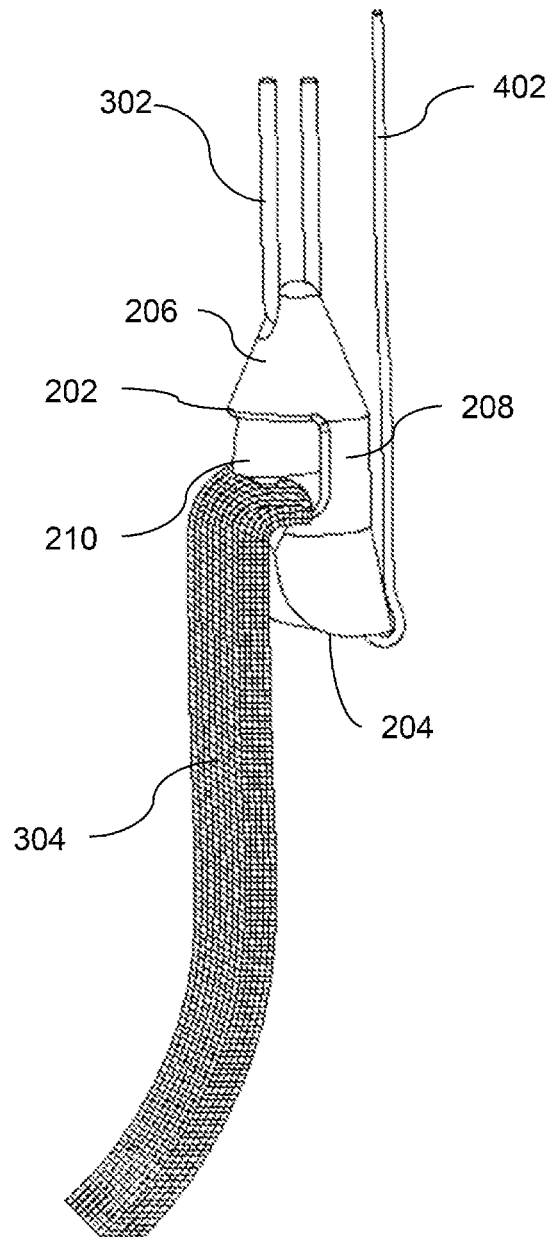


Fig. 4

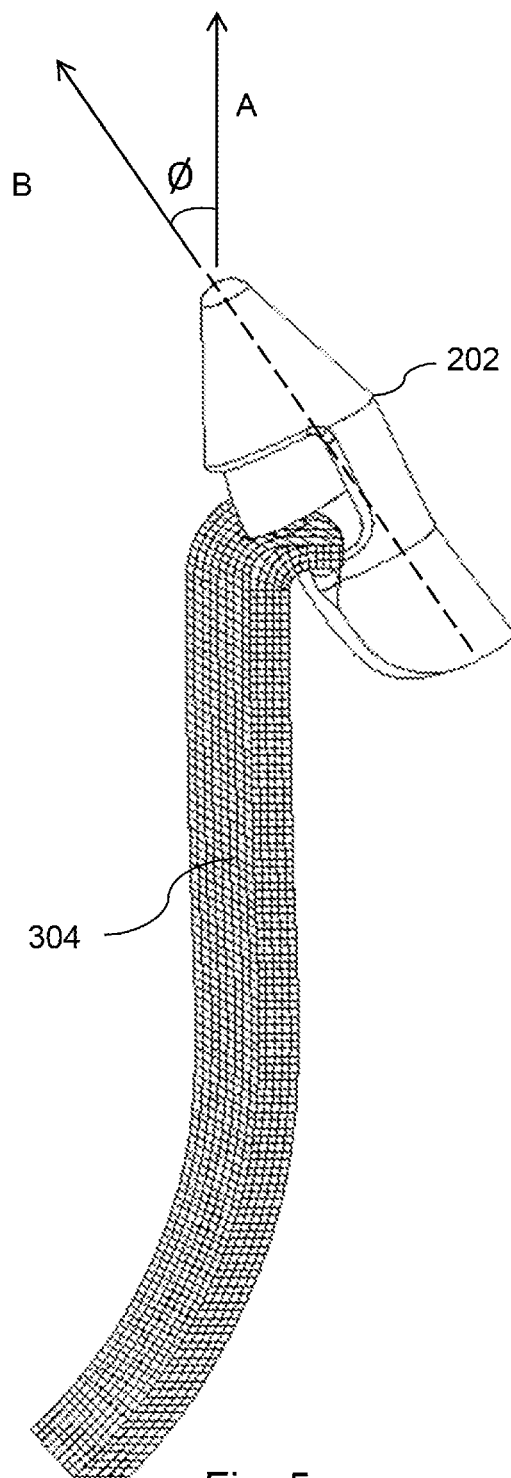


Fig. 5

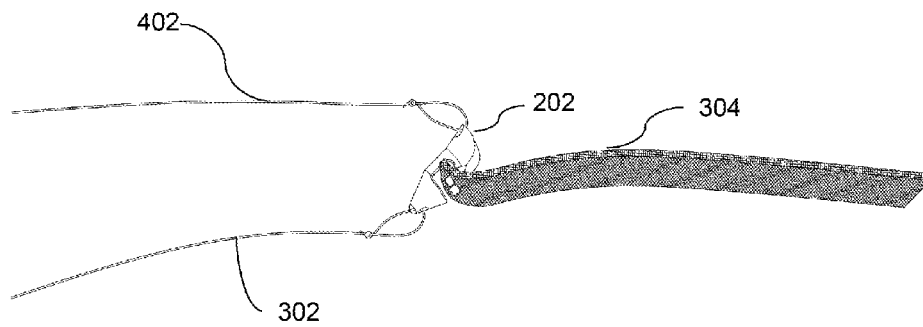


Fig. 6

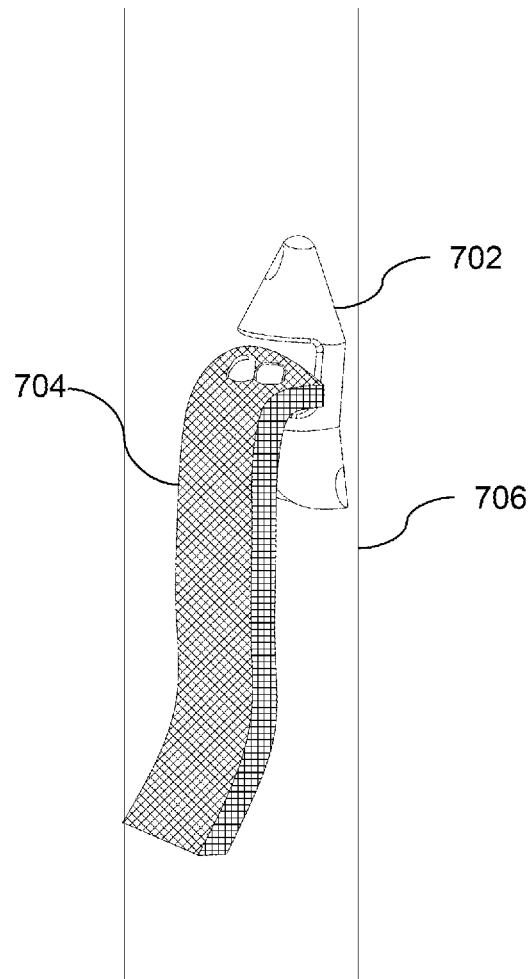


Fig. 7A

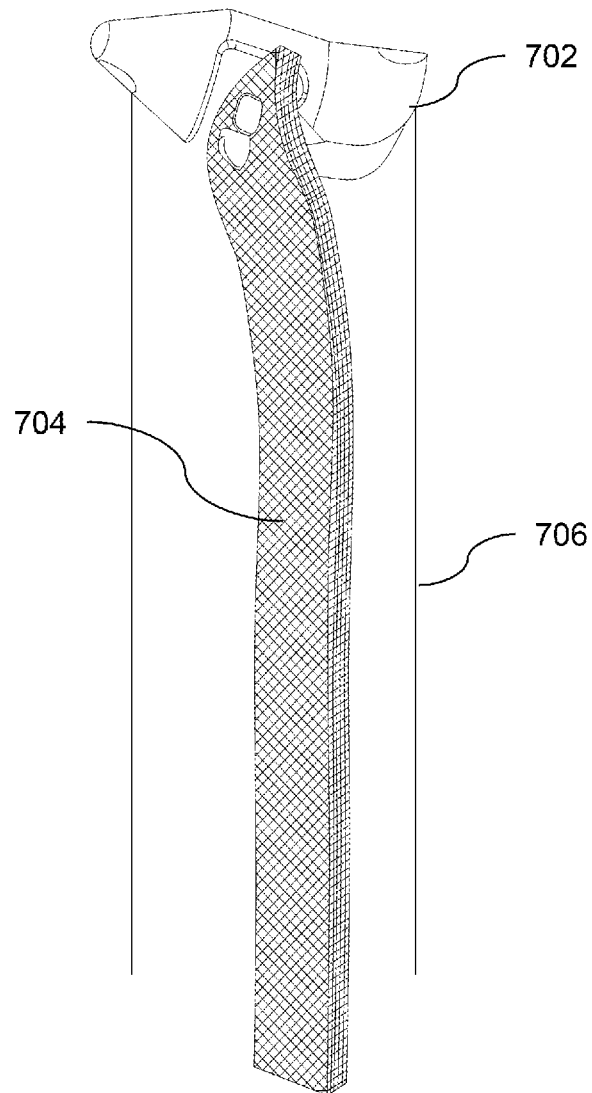


Fig. 7B

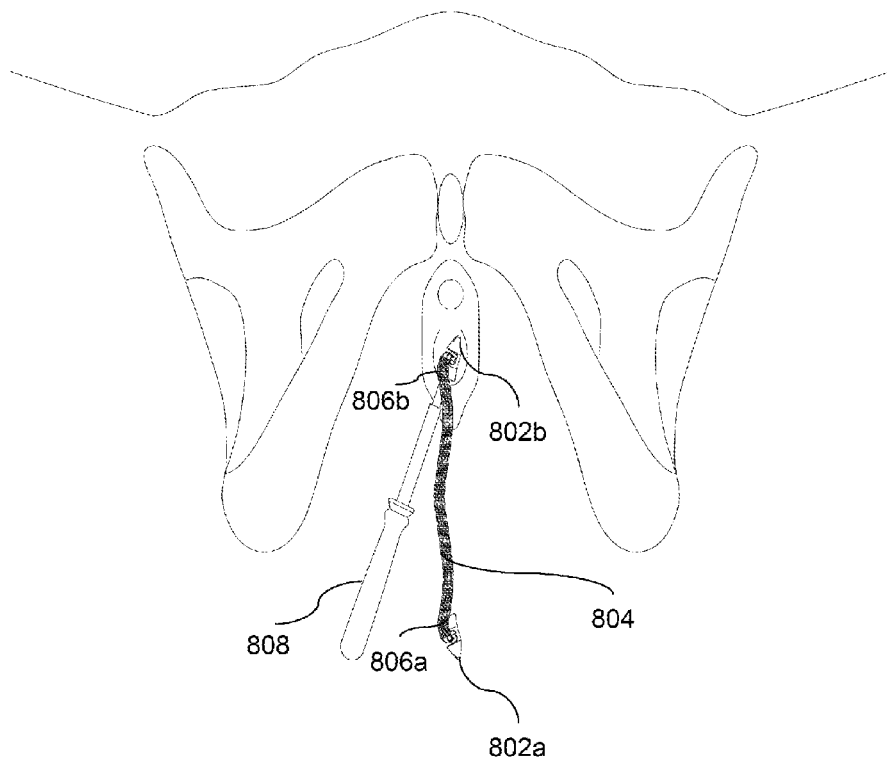


Fig. 8A

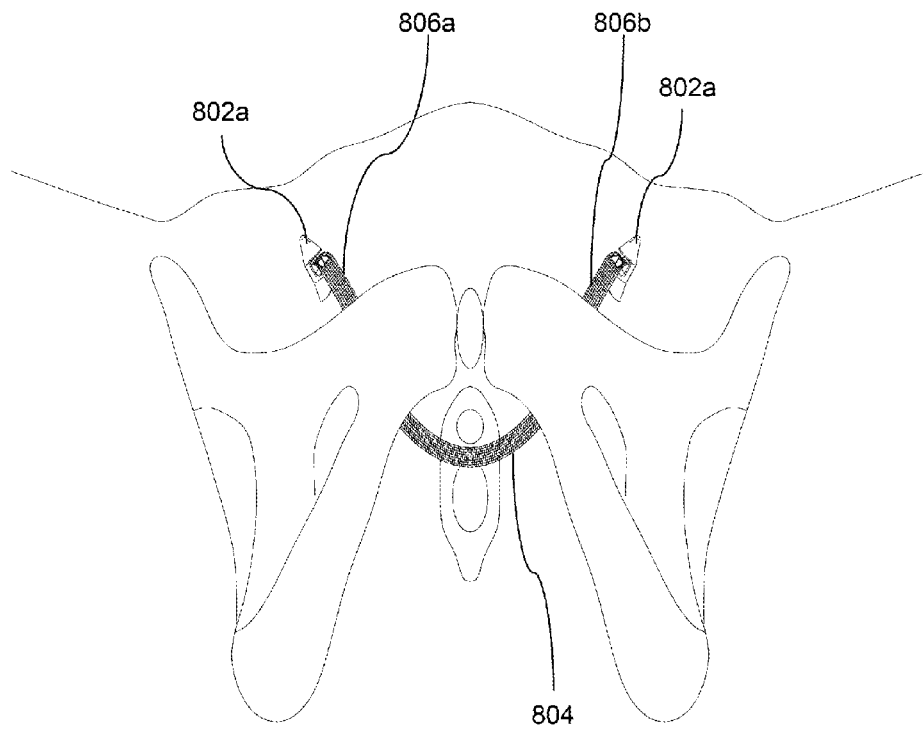


Fig. 8B

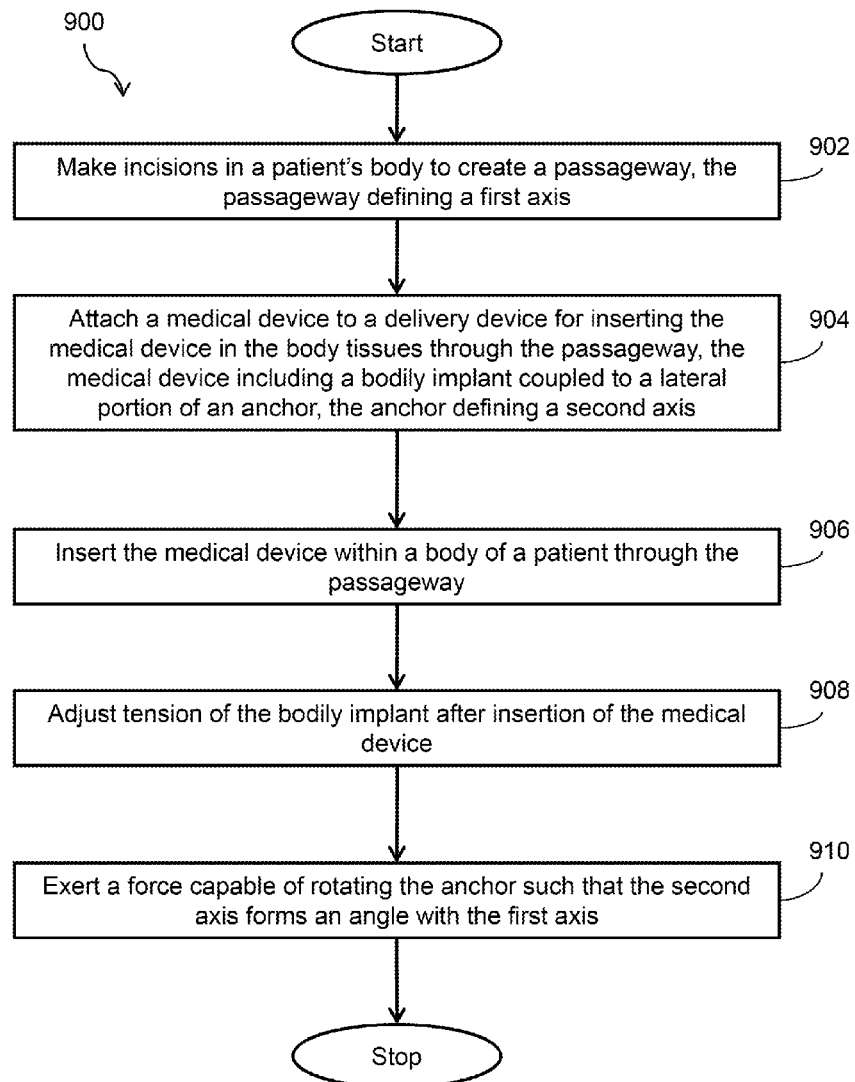


Fig. 9

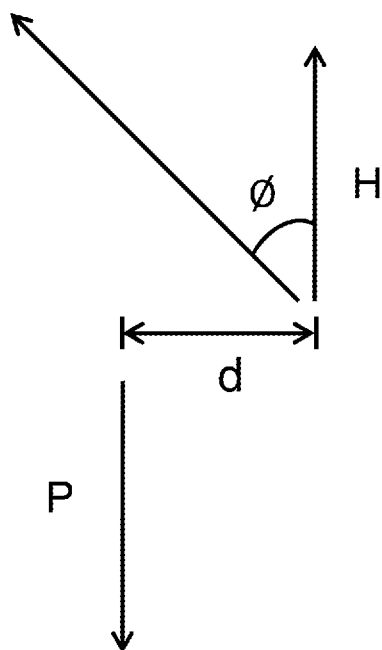


Fig. 10

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ANCHORS FOR BODILY IMPLANTS AND METHODS FOR ANCHORING BODILY IMPLANTS INTO A PATIENT'S BODY

CROSS-REFERENCE TO RELATED APPLICATION

This application is a Nonprovisional of, and claims priority to, U.S. patent application Ser. No. 61/485,388, filed May 12, 2011, entitled "ANCHORS FOR BODILY IMPLANTS AND METHODS FOR ANCHORING BODILY IMPLANTS INTO A PATIENT'S BODY", which is incorporated by reference herein in its entirety.

BACKGROUND

1. Field

The invention generally relates to medical devices and procedures, and more particularly to anchors for bodily implants and methods for anchoring the bodily implants into a patient's body.

2. Description of the Related Art

A common practice while inserting bodily implants, such as slings used in the treatment of urinary incontinence or fecal incontinence, is to use anchors. An anchor assists in holding a bodily implant and prevents it from being dislodged from its intended location with respect to an anatomy of a patient's body. The anchor works by engaging with surrounding anatomy and creating sufficient force to hold the bodily implant in its intended position.

Existing anchors are designed with anchoring protrusions. These protrusions vary in size based on the holding force required to anchor the bodily implants. In some existing anchors, the greater the holding force required, the longer the protrusions are. In some existing anchors, the protrusions are sharpened at their distal ends to ensure engagement of the anchors with surrounding tissues within the patient's body. However, it may be undesirable to leave the bodily implants, which have sharp edges protruding outwards, within the patient's body as the sharp edges may damage the surrounding tissues causing pain and discomfort. Further, such anchors may also cause damage to internal tissues of the patient's body during insertion and removal of anchors.

Thus, there is a need for an anchor that precludes the need for protrusions with sharp edges. Further, there is a need for an anchor that can exert a holding force on a bodily implant to anchor it at a suitable location in the patient's body.

SUMMARY

An anchor is provided for anchoring a bodily implant within a body of a patient. The anchor includes an implant engaging portion for engaging the bodily implant, wherein the implant engaging portion is disposed on a lateral portion of the anchor. The anchor further includes a distal end portion configured to pass through a passageway in the patient's body, the passageway defining a first axis and a proximal end portion disposed longitudinally opposite to the distal end portion on the anchor. The anchor defines a second axis extending from the distal end portion to the proximal end portion. The anchor is configured to rotate when a force is applied to the bodily implant such that the second axis defined by the anchor forms an angle with the first axis defined by the passageway.

BRIEF DESCRIPTION OF THE FIGURES

The invention and the following detailed description of certain embodiments thereof may be understood with reference to the following figures:

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FIG. 1 is a schematic diagram of an anchor affixed to an end portion of a bodily implant, in accordance with an embodiment of the present invention.

FIGS. 2A-2D illustrate perspective views of an anchor for affixing an end portion of a bodily implant, in accordance with various embodiments of the present invention.

FIG. 3 is a perspective view of an anchor with a delivery lead coupled to a distal end portion of the anchor, in accordance with an embodiment of the present invention.

FIG. 4 is a perspective view of an anchor with a tilt control lead coupled to a proximal end portion of an anchor, in accordance with an embodiment of the present invention.

FIG. 5 is a perspective view of an anchor in a rotated configuration depicting an angle formed between a first axis and a second axis.

FIG. 6 is a perspective view of an anchor coupled to a delivery lead and a tilt control lead, in accordance with another embodiment of the present invention.

FIG. 7A illustrates an exploded perspective view of an anchor within a bodily passageway during delivery.

FIG. 7B illustrates an exploded perspective view of an anchor within a bodily passageway after being rotated by an angle.

FIGS. 8A and 8B depict an illustrative method of implanting a bodily implant in a periurethral tissue of a patient, in accordance with an embodiment of the present invention.

FIG. 9 is a flowchart illustrating a method of implanting a bodily implant in a body of a patient, in accordance with an embodiment of the present invention.

FIG. 10 is a schematic diagram illustrating the mechanics of the forces, in accordance with an embodiment of the present invention.

DETAILED DESCRIPTION

Detailed embodiments of the present invention are disclosed herein; however, it is to be understood that the disclosed embodiments are merely exemplary of the invention, which may be embodied in various forms. Therefore, specific structural and functional details disclosed herein are not to be interpreted as limiting, but merely as a basis for the claims and as a representative basis for teaching one skilled in the art to variously employ the present invention in virtually any appropriately detailed structure. Further, the terms and phrases used herein are not intended to be limiting, but rather to provide an understandable description of the invention.

The terms "a" or "an," as used herein, are defined as one or more than one. The term "another," as used herein, is defined as at least a second or more. The terms "including" and/or "having", as used herein, are defined as comprising (i.e., open transition). The term "coupled" or "operatively coupled," as used herein, is defined as connected, although not necessarily directly and mechanically.

In general, the invention is directed to systems, methods, and devices for treating urinary incontinence. As described below in various illustrative embodiments, the invention provides systems, methods, and devices employing an improved soft tissue anchor termed as anchor for anchoring an end of a bodily implant in place, at least temporarily within a body of a patient. In accordance with several other embodiments, the invention may be used for the treatment of fecal incontinence as well.

The term patient may be used for a person who benefits from the anchors disclosed in the present invention. For example, the patient can be a person whose body receives the bodily implant with the disclosed anchor at its end in a sur-

gical treatment. For example, in some embodiments, the patient may be a human female, a human male, or any other mammal.

The terms proximal and distal described in relation to various devices, apparatuses, and components as discussed in the subsequent text of the present invention are referred with a point of reference. The point of reference, as used in this description, is a perspective of an operator. The operator may be a surgeon, a physician, a nurse, a doctor, a technician, and the like who may perform the procedure of delivery and placement of the bodily implants into the patient's body as described in the present invention. The term proximal refers to an area or portion that is closer or closest to the operator during a placement procedure. The term distal refers to an area or portion that is further or farthest from the operator.

FIG. 1 is a schematic diagram of an anchor **102** affixed to an end portion of a bodily implant **104**, in accordance with an embodiment of the present invention. The anchor **102** and the bodily implant **104** that are configured to be placed inside a patient's body can together be hereafter referred to as a medical device **100** for the simplicity of the description.

The anchor **102** includes a proximal end portion **106**, a distal end portion **108**, and a medial portion **110**. The distal end portion **108** is configured to pass through a passageway in the patient's body. For example, in some embodiments, the distal end portion **108** is configured to pass through a passageway that is formed during insertion of the anchor **102**. In some embodiments, the passageway defines a first axis A. The proximal end portion **106** is disposed longitudinally opposite to the distal end portion **108** on the anchor **102** such that the proximal end portion **106** and the distal end portion **108** lie at two ends of the anchor **102**. The medial portion **110** (that, in some embodiments, integrally forms a middle part of the anchor **102** between the proximal end portion **106** and the distal end portion **108**) further includes an implant engaging portion **112** for engaging the bodily implant **104**. The implant engaging portion **112** is disposed on a lateral or side portion of the anchor **102** such that the bodily implant **104** is asymmetrically attached to the anchor **102**. Further, the implant engaging portion **112** extends from the proximal end portion **106** toward the distal end portion **108** such that the longitudinal axis of the implant engaging portion **112** is parallel to the longitudinal axis of the anchor **102**.

The bodily implant **104** can be coupled to the implant engaging portion **112** in various ways. There can be different types of mechanisms to couple the bodily implant, in accordance with various embodiments. For example, the bodily implant **104** can be pierced within the implant engaging portion **112** and subsequently glued, stapled, or tied to the implant engaging portion **112**. Numerous types of designs of the implant engaging portion **112** are possible depending on the nature of mechanism of engaging the bodily implant **104**.

In accordance with some embodiments, the implant engaging portion **112** includes at least one protuberance for engaging the bodily implant **104** therewith. The at least one protuberance is designed to extend longitudinally from a lower portion of the implant engaging portion **112** and configured to hold the bodily implant **104** at place. The at least one protuberance may be a small extension or projection extending from the lateral or side portion of the anchor **102**.

In some embodiments, there can be only one protuberance. In some other embodiments, there can be two protuberances, a first protuberance and a second protuberance. The first and the second protuberances are configured so that the bodily implant **104** can be pierced through them and fixed therein. In some embodiments, the first protuberance and the second protuberance are configured to interlock with each other and

engage the bodily implant **104** therein. In accordance with these embodiments, the first protuberance may include a male coupling member and the second protuberance may include a female coupling member such that the coupling members may fit in an interlocked manner.

In some embodiments, the at least one protuberance may include a movable locking mechanism for engaging the bodily implant **104**. The movable locking mechanism may be configured to latch or lock the bodily implant **104** to fixedly couple the implant **104** to the engaging portion **112**. In some embodiments, the movable locking mechanism may be operated through a sliding mechanism such that the bodily implant **104** is latched or coupled to the implant engaging portion **112** by slidably moving the at least one protuberance relative to the anchor **102**. At least one opening may be provided on the implant engaging portion **112** such that the at least one protuberance may slidably fit into the at least one opening.

In still various other embodiments, several types of locking, latching, and engaging mechanisms may be provided on the implant engaging portion **112** that are capable of holding and engaging the bodily implant **104**.

In embodiments, the anchor **102** is elongated in nature such that a length of the anchor **102** which extends longitudinally is substantially more than a width of the anchor **102** which extends transversely. The anchor **102** defines an axis (second axis B) extending from the distal end portion **108** toward the proximal end portion **106**.

The proximal end portion **106** and the distal end portion **108** may have any suitable size and shape. In some embodiments, the distal end portion **108** is substantially conical. In other embodiments, the distal end portion **108** may be substantially rectangular, circular, and the like. In some embodiments, the proximal end portion **106** is substantially cylindrical. In other embodiments, the proximal end portion **106** is substantially rectangular, circular, and the like. A tip portion of the distal end portion **108** that is configured to pass through the passageway in the patient's body may be shaped conically and sharp in nature. In general, the anchor **102** may have any shape and size that is suitable for affixing the anchor **102** within an anatomical membrane, muscle, ligament, soft tissue, bone or any other anatomical site.

In some embodiments, the anchor **102** may be made of any suitable biocompatible material. In other embodiments, the anchor **102** may be made, for example, of a synthetic material such as nylon, polyethylene, polyester, polypropylene, fluoropolymers or a co-polymer thereof. In some other embodiments, they may be formed, at least in part, from a mammalian tissue material such as bovine, porcine, equine, human cadaveric or engineered tissue. In still other embodiments, the material of the anchor **102** may include a combination of synthetic and mammalian tissue/biocompatible materials. In some embodiments, the anchor **102** is made of a metal, ceramic, polymer, magnet, or an alloy.

According to some embodiments, at least a portion of the anchor **102** is biodegradable and may also dissolve and/or be absorbed by the patient's tissues. Exemplary biodegradable materials that may be employed for at least a portion of the anchor **102** include, but are not limited to, polylactic acid, polyglycolic acid, and copolymers and mixtures thereof, such as poly(L-lactide) (PLLA), poly(D,L-lactide) (PLA), polyglycolic acid [polyglycolide (PGA)], poly(L-lactide-co-D,L-lactide) (PLLA/PLA), poly(L-lactide-co-glycolide) (PLLA/PGA), poly(D,L-lactide-co-glycolide) (PLA/PGA), poly(glycolide-co-trimethylene carbonate) (PGA/PTMC), poly(D,L-lactide-co-caprolactone) (PLA/PCL), and poly(glycolide-co-caprolactone) (PGA/PCL); polyethylene oxide

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(PEO); polydioxanone (PDS); polypropylene fumarate; polydepsipeptides, poly(ethyl glutamate-co-glutamic acid), poly(tert-butyloxy-carbonylmethyl glutamate); polycaprolactone (PCL), poly(hydroxy butyrate), polycaprolactone co-butylacrylate, polyhydroxybutyrate (PHBT) and copolymers of polyhydroxybutyrate; polyphosphazenes, polyphosphate ester); maleic anhydride copolymers, polyiminocarbonates, poly[(97.5% dimethyl-trimethylene carbonate)-co-(2.5% trimethylene carbonate)], cyanoacrylate, hydroxypropylmethylcellulose; polysaccharides, such as hyaluronic acid, chitosan and regenerate cellulose; poly(amino acid) and proteins, such as gelatin and collagen; and mixtures and copolymers thereof.

In some embodiments, the anchor **102** may be configured to be dissolved within a particular time range. The anchor **102** may be configured, for example, to substantially absorb (or have a portion that substantially absorbs) into the patient's tissues within about 2, 4, 6 or 8 or more weeks from the time the bodily implant **104** is implanted. Preferably, the anchor **102** remain structurally intact long enough for scar tissue and/or other neighboring cells or tissues to grow into the bodily implant **104** to effectively anchor it in place.

The bodily implant **104** that is affixed with the use of the anchor **102** is configured to be placed within the patient's body and support a portion of the body. For example, the bodily implant **104** can be shaped and sized to support a portion of the body around a bladder, urethra, anal canal, rectum, and anus of the patient. The bodily implant **104** has a first end portion **114** and a second end portion **116** such that the bodily implant **104** extends along a length between the first end portion **114** and the second end portion **116**. The length and width of the bodily implant **104** may vary based on its intended use. The bodily implant **104** can be of a variety of sizes, shapes, and configurations depending on the intended use and locations of placement of the bodily implant **104**.

In some embodiments, the bodily implant **104** is formed of a material that allows tissue in-growth after implantation. Various types of woven tapes, fabrics, or meshes may be utilized in the fabrication and manufacturing of the bodily implant **104**, in accordance with various embodiments of the present invention. The bodily implant **104** may utilize a variety of mesh materials and may be designed in a variety of forms. An example of a mesh utilized in the bodily implant **104** is Polyform® Synthetic Mesh developed by the Boston Scientific Corporation. The Polyform® Synthetic Mesh is made from uncoated monofilament macro-porous polypropylene. The bodily implant **104** may also be made from a biological material or a cadaveric tissue. In some embodiments, the bodily implant **104** has a smooth surface. In such embodiments, the smooth surface may avoid or reduce irritation on adjacent body tissues during mesh-tissue interactions. Additionally, the bodily implant **104** may be stretchable and flexible to adapt movements in accordance with the anatomy of the human body and reduce suture or anchor pullout. Furthermore, softness, lightness, conformity, and strength are certain other attributes required in the bodily implant **104** for efficient tissue repair and implantation. In an embodiment, the bodily implant **104** can have a coating. For example, the bodily implant **104** can be coated with an antimicrobial agent and/or an antifungal agent.

FIG. 2A-2D illustrate perspective views of an anchor **202** for affixing an end portion of a bodily implant such as the bodily implant **104** as illustrated in FIG. 1, in accordance with an embodiment of the present invention. As depicted, the anchor **202** includes a proximal end portion **204**, a distal end portion **206** and a medial portion **208**.

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The distal end portion **206** is configured to pass through a passageway in the patient's body such that the passageway defines a first axis A. In some embodiments, the distal end portion **206** is configured to create the passageway as it is inserted into the body of the patient. The proximal end portion **204** is disposed longitudinally opposite to the distal end portion **206** on the anchor **202**. The medial portion **208** further includes an implant engaging portion **210** for engaging the bodily implant. The implant engaging portion **210** is disposed on a lateral or side portion of the anchor **202** such that the bodily implant is asymmetrically attached to the anchor **202**. The implant engaging portion **210** extends axially from the proximal end portion **204** toward the distal end portion **206** and positioned asymmetrically at a lateral part of the medial portion **208**. The term asymmetric attachment means that the bodily implant is coupled at only one side of the longitudinal axis of the anchor **202** and not on both sides. The effect of asymmetric attachment is available along only one lateral side rather than on the longitudinal axis at the center of the anchor **202**.

According to some embodiments, the implant engaging portion **210** is formed integrally with the medial portion of the anchor **208**. In other embodiments, the implant engaging portion **210** is separable from the medial portion **208** such that it is configured to removably fit into the medial portion **208**. The anchor **202** defines a second axis B extending from the distal end portion **206** toward the proximal end portion **204**. The second axis B coincides with the longitudinal axis along the length of the anchor **202**.

As illustrated in FIGS. 2A-2D, the implant engaging portion **210** further includes two protuberances, a first protuberance **212a** and a second protuberance **212b**. There can be different types of mechanisms to couple the bodily implant through the first protuberance **212a** and the second protuberance **212b**, in accordance with various embodiments. For example, the bodily implant can be pierced through the protuberances **212a** and **212b**, and subsequently glued, stapled, or tied. Numerous types of designs of the protuberances **212a** and **212b** are possible depending on the nature of mechanism for engaging the bodily implant.

In some embodiments, the first protuberance **212a** and the second protuberance **212b** are designed to extend longitudinally from a lower portion of the implant engaging portion **210** and configured to hold the bodily implant at place. The protuberances **212a** and **212b** may be designed in the form of small extensions or projections extending from the lateral or side portion of the anchor **202**, as illustrated in FIG. 2A.

In some embodiments, the first protuberance **212a** and the second protuberance **212b** are configured to interlock with each other and engage the bodily implant therein. In accordance with these embodiments, the first protuberance **212a** may include a female coupling member **216** and the second protuberance **212b** may include a male coupling member **218**, as shown in FIG. 2B. The coupling members **216** and **218** can fit into one another for interlocking.

In some embodiments, the first protuberance **212a** and the second protuberance **212b** may include a movable locking mechanism **220** for engaging the bodily implant, as shown in FIGS. 2C and 2D. The movable locking mechanism **220** may be configured to latch or lock the bodily implant to fixedly couple the implant to the engaging portion **210**. In some embodiments, the movable locking mechanism **220** may be operated through a sliding mechanism such that the bodily implant is latched or coupled to the implant engaging portion by slidably moving the protuberances **212a** and **212b** relative to the anchor **102**. FIG. 2D shows a latched configuration achieved after sliding. A latch **220** may be provided with the

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sliding mechanism **220** to retain the protuberances **212a** and **212b** in the latched configuration. Two openings (not shown) may be provided on the implant engaging portion **210** such that the protuberances **212a** and **212b** may slidably fit into the openings.

In still various other embodiments, several types of locking, latching, and engaging mechanisms may be provided that are capable of locking and latching the body implant with the protuberances **212a** and **212b**. In accordance with various embodiments, the first protuberance **212a** and the second protuberance **212b** are designed to be projectionless and barbless such that these protuberances, specifically their tip portions do not harm and irritate the body tissues.

The anchor **202** can have a variety of shapes and sizes similar to the anchor **102** as described in conjunction with FIG. 1. Similarly, the material and composition of the anchor **202** can vary as described in conjunction with FIG. 1.

As illustrated in FIGS. 2A-2D, the anchor **202** includes or defines a first opening or lumen **214** defined on the distal end portion **206** of the anchor **202**. The first opening **214** is disposed on the distal end portion **206** such that the lateral ends of the first opening **214** passes through two conical edges of the distal end portion **206**. A second opening or lumen (not shown) is provided on the proximal end portion **204**. The second opening can be provided at a lateral surface of the proximal end portion **204** or at a bottom surface of the anchor **202**. In some embodiments, the second opening is asymmetrically disposed on the anchor **202** such that a distance of the second opening from a first lateral edge of the anchor **202** is more than a distance of the second opening from a second lateral edge of the anchor **202**. In some embodiments, the second opening may extend from a lateral edge of the anchor **202** to a bottom portion of the anchor **202**. In accordance with various embodiments, the shape and size of the first and the second openings (hereafter referred to as openings together) may vary based on the intended use and the requirements.

The first opening **214** is defined to receive and engage a first lead termed as a delivery lead **302** with the anchor **202**, as illustrated in FIG. 3. In some embodiments, the delivery lead can include a loop at one of its end portions configured to be coupled to a delivery tool such as a surgical needle. In some embodiments, the loop may be some kind of a suture loop. The delivery lead **302** can be brought through body tissues to assist in the delivery of a bodily implant such as the bodily implant **304**. The anchor **202** can be forced into the passageway within the body tissues by applying forces on the delivery lead **302**. As illustrated in FIG. 4, the second opening is defined to receive and engage a second lead termed as a tilt control lead **402**. The tilt control lead **402** can be brought through the body tissues along with the delivery lead **302**. The tilt control lead **402** is used to rotate the anchor **202** such that an angle θ is formed between the first axis A (defined by the passageway) and the second axis B (defined by the anchor) in FIG. 5. The angle θ thus formed assists in anchoring the bodily implant such as the bodily implant **304** with the body tissues. FIG. 6 illustrates a perspective view of the delivery lead **302** and the tilt control lead **402** coupled to the anchor **202**. As shown, the leads **302** and **402** can form a loop at the coupling end such that the leads **302** and **402** can be removed easily after placement of the anchor **202** by cutting the loops.

FIG. 7A illustrates an exploded perspective view of an anchor **702** within a bodily passageway **706** during delivery. The anchor **702** is coupled to a bodily implant **704**. The longitudinal axis of the anchor **702** is substantially parallel to the direction of the bodily passageway **706** during delivery. FIG. 7B illustrates an exploded view of the anchor **702** after being rotated by an angle upon placement. In this configura-

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tion, the anchor **702** engages with the bodily tissues since the length of the anchor **702** (along its longitudinal axis) is more than its width. The measure of the angle by which the anchor **702** is rotated can vary depending on the amount of rotational torque or turning momentum generated to cause the rotation of the anchor **702**.

FIGS. 8A and 8B depict an illustrative method of implanting a bodily implant such as the bodily implant **804** in a periurethral tissue of a patient to form a platform under a urethra of the patient. As illustrated in FIG. 8A, a first anchor **802a** is fitted on a first end portion **806a** of the bodily implant **804** and a second anchor **802b** is fitted on a second end portion **806b** of the bodily implant **804**. The two anchors **802a** and **802b** are configured to anchor and hold the bodily implant **804** at two end portions **806a** and **806b** such that the bodily implant **804** is fixedly supported inside the body tissues in an appropriate tension. In some embodiments, at least one of the anchors **806a** and **806b** are similar to the anchor **102** illustrated in conjunction with FIG. 1. In other embodiments, at least one of the anchors **806a** and **806b** are similar to the anchor **202** illustrated in conjunction with FIG. 2.

FIG. 9 is a flowchart illustrating a method **900** for anchoring a bodily implant such as the bodily implant **804** within a body of a patient. In accordance with various embodiments, an anchor such as the anchor **802a** can be used to fix the bodily implant **804** in place. The anchor **802a** has a length that is substantially more than its width. During delivery, the anchor **802a** is inserted along a longitudinal direction, which is parallel to axis of the bodily passageway. After insertion, the anchor **802a** is rotated by an angle with respect to the bodily passageway. In this configuration, the longitudinal direction of the anchor **802a** is substantially perpendicular to the passageway. Since the length of the anchor **802a** is more than the width of the bodily passageway, the anchor **802a** gets engaged within the bodily tissues. Similarly, the anchor **802b** can also be used to fix the other end of the bodily implant **804**. The configuration of the anchor (during delivery and after rotation) is illustrated in FIGS. 7A and 7B, respectively.

Referring now to FIGS. 8A, 8B, and 9 together, a specific method for implanting and anchoring a bodily implant such as the bodily implant **804** is described in accordance with an embodiment of the present invention. At step **902**, an incision is made in an anterior vaginal wall and dissected bilaterally to the interior portion of an inferior pubic ramus of the patient. The vaginal incision creates a passageway from the vaginal opening to urethral sphincter that is responsible for controlling the flow of urine. The vaginal incision allows the bodily implant **804** to be placed correctly under the urethra, without passing a delivery tool **808** through the retropubic space and abdominal wall unknowingly causing damage. The direction of the passageway defines a first axis. The first axis is along the direction of insertion through the body tissues. In some embodiments, an operator further makes second and third incisions in groin areas—one on a left groin area and the other on a right groin area on either side of the pubis. In other embodiments, the second and the third incisions can be made in the obturator membrane or in the abdomen.

At step **904**, the operator attaches/couples a medical device to the delivery tool **808**. The medical device includes the bodily implant **804** coupled to lateral portions of the anchors **802a** and **802b** at its two end portions **806a** and **806b** as illustrated in FIGS. 8A and 8B. For example, a first anchor **802a** is coupled to a first end portion **806a** of the bodily implant **804** through a first implant engaging portion and a second anchor **802b** is coupled to a second end portion **806b** of the bodily implant **804** through a second implant engaging portion. The first anchor **806a** defines a second axis extending

from a distal end portion toward a proximal end portion of the first anchor **806a**. The second axis for the first anchor **806a** coincides with the longitudinal axis along the length of the first anchor **806a**. Similarly, the anchor **806b** also defines a second axis extending from a distal end portion toward a proximal end portion of the second anchor **806b**. The second axis for the second anchor **806b** coincides with the longitudinal axis along the length of the second anchor **806b**.

The delivery tool **808** may be an elongated member such as a surgical needle that may be fitted to an anchor such as the anchor **806a** and **806b** during delivery of a bodily implant such as the bodily implant **804** as shown in FIG. **8A**. The delivery tool **808** may include a shaft that may be substantially straight, curved or include both curved and straight portions. In some embodiments, a distal tip of the shaft is conically shaped to provide a sharp end facilitating insertion of the bodily implant **804** and the anchors **806a** and **806b** inside the body tissues.

At step **906**, the medical device is inserted through the passageway in a patient's body. In some embodiments, the delivery tool **808** carrying the medical device is inserted through the vaginal incision that acts as the passageway for advancing the delivery tool **808**. A force of insertion applied by the operator moves the medical device within the patient's body. Fingers of the operator may guide the delivery tool **808** inside the body to avoid blind delivery and hence, achieve effective advancement inside the body. The maximum depth of advancement through the vaginal incision must be limited to avoid perforation of the bladder wall.

In some embodiments, the anchors **802a** and **802b** of the medical device can be directly coupled to the delivery tool **808** for insertion into the body. The anchors **806a** and **806b** can include slots or interfaces disposed on their proximal end portions such that a distal tip portion or a needle tip of the delivery tool **808** can be engaged through the slots or interfaces of the anchors **806a** and **806b**. This provides a coupling of the delivery tool **808** with the anchors **806a** and **806b** such that an engagement of the anchors **806a** and **806b** with the delivery tool **808** through the slots or the interfaces ensure proper delivery and insertion of the medical device into the body tissues.

In accordance with some other embodiments, a delivery lead such as the delivery lead **302** may be utilized for inserting the medical device into the body. In some embodiments, the delivery lead **302** includes a loop that can be coupled to the delivery tool **808**. The loop is configured to be hooked to the delivery tool such that the delivery lead **302** is pushed into the body, upon insertion of the delivery tool **808** through the vaginal incision, and comes out through groin area or abdomen of the patient. The delivery lead **302** can be finally unhooked from the tool **808** and the tool **808** is pulled out through the vaginal incision backward. In some embodiments, the delivery lead **302** is then pulled outside to leave the anchors **802a** and **802b** inside the body. In accordance with various other embodiments, several other types of bodily incisions and insertion mechanisms may be employed to insert the medical device inside the patient's body depending on the preference of an operator or a physician and the condition of the patient to be treated.

Once the medical device is inserted and placed within the body, the tension of the bodily implant **804** is adjusted at step **808**. In some embodiments, the operator may adjust the tension of the bodily implant **804** by stretching it manually after placement at its targeted location. In other embodiments, the tension may be adjusted by a tension member such as a suture. Various other procedures of adjusting tension may be utilized without limitations.

After an appropriate tension is confirmed in the bodily implant **804**, it is anchored within the body tissues at step **810** by exerting a force capable of rotating the anchors **802a** and **802b** such that the second axis of the anchors **802a** and **802b** forms an angle with the first axis with respect to the anchors **802a** and **802b**. The anchoring is done by using the two anchors **802a** and **802b** that are coupled at the two end portions **806a** and **806b** of the bodily implant **804** through their implant engaging portions. For example, the first end portion **806a** of the bodily implant **804** is anchored in a first portion of the body tissues using the first anchor **802a** and the second end portion **806b** of the bodily implant **804** is anchored in a second portion of the body tissues using the second anchor **802b**.

In some embodiments, the anchoring of the bodily implant **804** is done by first exerting a force on the bodily implant **804** outward at a portion that extends and hangs out of the patient's body. For example, a force may be exerted on the bodily implant **804** outward to anchor the first end portion **806a** of the bodily implant **804** using the first anchor **802a**. This causes the development of a pulling force that acts in a direction opposite to the direction of the insertion as a result of an interaction of the bodily implant **804** with the body tissues. Since the first end portion **806a** of the bodily implant **804** is asymmetrically coupled on the lateral portion of the first anchor **802a** within the implant engaging portion, the pulling force develops at the lateral portion of the first anchor **802a** eccentrically and not to the centre.

As a result of the development of the pulling force, a holding force starts developing at a catching point. The catching point is present on the bottom proximal right side of the anchor **802a**. When the anchor **802a** is placed inside the body and the implant is pulled down, the anchor **802a** slightly goes down and hits the passageway at the catching point. This helps pivot the anchor **802a** into a rotated configuration with respect to the passageway. The holding force at the catching point and the pulling force along the bodily implant **804** form a force couple or a turning momentum, which rotates the first anchor **802a**. The rotation of the anchor **802a** makes the distance between the forces of the turning momentum greater, thereby increasing the turning momentum even more. In this scenario, snow cone effect develops that tilts the anchor **802a** by an angle formed between the first axis and the second axis. Therefore, the first anchor **802a** is lodged within the tissues and creates a large holding force capable of restoring the anchor **802a** at the desired position. The mechanics of the forces is illustrated in FIG. **10**. As illustrated in the FIG. **10**, 'P' represents the pulling force applied along the bodily implant **804**, 'H' represents the holding force generated as a result of the pulling force, 'Ø' represents the angle formed between the first axis and the second axis after the first anchor **802a** rotates, 'd' represents a distance between the lines of action of the two forces—the holding force and the pulling force.

The turning momentum at an engagement point of the implant engaging portion and the bodily implant **804** that causes rotation of the anchor **802a** tries to bring it downward with respect to the body tissues. This changes the direction of the first anchor **802a** and its proximal end portion now faces opposite to the lateral edge of the bodily implant **804** in a transverse direction, thereby engaging it with the body tissues at the catching point. The angle of rotation resulting from the effect of the turning momentum may be measured as an angle formed between the first axis defined by the passageway and the second axis defined by the longitudinal direction of the first anchor **802a** joining the proximal and distal end portions

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of the first anchor **802a**. The angle thus formed between the first axis and the second axis as a result of rotation is depicted in FIGS. **5** and **10** as \emptyset .

The anchoring of the first end portion **806a** of the bodily implant **804** is achieved by rotating the anchor **802a** on application of a pulling force on the bodily implant **804** as described above. In accordance with other embodiments, the turning momentum that is capable of rotating the first anchor **802a** can be generated by pulling a second lead termed as a tilt control **402** as illustrated in FIG. **4** from its distal end. A proximal end of the tilt control lead **402** is coupled to the proximal end of the first anchor **802a** in a manner as described in conjunction with FIG. **4**. The operator may exert a force on the tilt control lead **402** to rotate the anchor **802a** such that the second axis defined by the longitudinal direction of the first anchor **802a** makes an angle \emptyset with the first axis defined by the passageway. In accordance with still other embodiments, a pulling force on the bodily implant **804** as well as a pulling force on the tilt control lead **402** can be applied together to achieve a desired angle \emptyset between the first axis and the second axis such that the anchor **802a** is appropriately lodged in the body tissues.

In a manner similar to the anchoring of the first end portion **806a** of the bodily implant **804** with the use of the first anchor **802a** at the first implant engaging portion, the second end portion **806b** of the bodily implant **804** may also be anchored using the second implant engaging portion of the second anchor **802b**. The second implant engaging portion is coupled at the second end portion **806b** of the bodily implant **804**. In this scenario, an angle is formed between the first axis and the second axis with respect to the second anchor **802b**. In some embodiments, the angle formed between the first axis and the second while anchoring the first anchor **802a** is same as the angle formed between the first axis and the second while anchoring the second anchor **802b**. In other embodiments, the angle formed between the first axis and the second while anchoring the first anchor **802a** is different than the angle formed between the first axis and the second while anchoring the second anchor **802b**. In accordance with various embodiments, the rotation angle \emptyset formed between the first axis and the second axis may vary based on the requirements such as the intended use and placement location of the bodily implant **804**.

In accordance with various embodiments, re-positioning of the bodily implant **804** may be done in case the bodily implant **804** is found to be placed incorrectly. In order to reposition the bodily implant **804**, the operator may exert a force on the delivery lead **302** coupled to the distal end portion of the anchor such as the anchor **802a** and **802b**. An appropriate force on the delivery lead **302** aligns the second axis with the first axis such that the longitudinal direction of the anchor such as the anchor **802a** and **802b** coincides with the direction of the passageway. Thus, the anchor such as the anchor **802a** and **802b** are no more in a rotated configuration. The operator adjusts the placement of the anchor (**802a** and **802a**) and finally rotates them in accordance with various embodiments described above.

In some embodiments, the anchors **802a** and **802b** can be left to stay inside the body tissues. In some other embodiments, the anchors **802a** and **802b** can be removed from the patient's body. The anchors **802a** and **802b** can be removed by exerting a force on the tilt control lead such that the anchors **802a** and **802b** are rotated by 180 degree (with respect to the direction of the passageway) to align the first axis and the second axis. This makes the distal ends of the anchors **802a** and **802b** face toward the direction of the passageway such that a simple pull applied on the distal ends of

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the anchors **802a** and **802b** can remove them outside the patient's body. In some other embodiments, the anchors (**802a** and **802b**) can be removed even without rotating through the 180 degree angle. In accordance with these embodiments, a simple pull is required at the delivery lead **302** or at the delivery lead **302** and the bodily implant **804** together to straighten the anchors (**802a** and **802b**) such that the first axis coincides with the second axis. In this scenario, a pull of magnitude equivalent to rotate the anchors (**802a** and **802b**) by an angle \emptyset and in opposite direction can straighten the anchors (**802a** and **802b**). The anchors (**802a** and **802b**) can be easily removed from the body by pulling them outside manually in a backward direction once they are in straight configuration.

The method for implanting and anchoring a bodily implant using anchors is described in conjunction with the bodily implant **804** and the anchors **802a** and **802b** above. However, the anchors such as **102** and **202** can also be used to anchor the bodily implant in accordance with various other embodiments of the present invention. Similarly, the bodily implant **104**, **304**, and various other kinds of bodily implants as used conventionally may also be employed.

In one embodiment, an anchor for anchoring a bodily implant within a body of a patient includes a distal end portion configured to pass through a passageway in the patient's body, the passageway defining a first axis and a proximal end portion disposed longitudinally opposite to the distal end portion on the anchor. The anchor defines a second axis extending from the distal end portion to the proximal end portion. A medial portion having an implant engaging portion for engaging the bodily implant, the implant engaging portion disposed on a lateral portion of the anchor. The anchor is configured to rotate such that the second axis defined by the anchor forms an angle with the first axis defined by the passageway upon rotation.

In some embodiments, the implant engaging portion includes at least one protuberance for engaging the bodily implant therewith. In some embodiments, the implant engaging portion includes a movable locking mechanism for engaging the bodily implant. In some embodiments, the implant engaging portion includes a slidable locking mechanism for engaging the bodily implant. In some embodiments, the implant engaging portion includes a first protuberance and a second protuberance configured to interlock with each other and engage the bodily implant therewithin.

In some embodiments, the proximal end portion defines an opening for coupling a tilt control lead with the anchor. In some embodiments, the anchor is configured to be rotated when a force is exerted on the tilt control lead. In some embodiments, the anchor is rotated to align the first axis and the second axis for removal of the anchor such that the distal end portion faces the passageway.

In some embodiments, the proximal end portion is substantially cylindrical. In some embodiments, the distal end portion is substantially conical. In some embodiments, the distal end portion defines an opening for engaging a delivery lead with the anchor. In some embodiments, the anchor is composed of at least one of a bio-compatible material, plastic, polypropylene, metal, ceramic, polymer, magnet, and alloy.

In some embodiments, a medical device is configured to be inserted within a body of a patient. The medical device includes a bodily implant and an anchor. The anchor includes a distal end portion configured to pass through a passageway in the patient's body, the passageway defining a first axis and a proximal end portion disposed longitudinally opposite to the distal end portion on the anchor. The anchor defines a second axis extending from the distal end portion to the

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proximal end portion and a medial portion having an implant engaging portion for engaging the bodily implant. The implant engaging portion disposed on a lateral portion of the anchor. The anchor is configured to rotate such that the second axis defined by the anchor forms an angle with the first axis defined by the passageway upon rotation.

In some embodiments, the implant engaging portion includes at least one protuberance for engaging the bodily implant therewith. In some embodiments, the implant engaging portion includes a movable locking mechanism for engaging the bodily implant. In some embodiments, the implant engaging portion includes a slidable locking mechanism for engaging the bodily implant. In some embodiments, the implant engaging portion includes a first protuberance and a second protuberance configured to interlock with each other and engage the bodily implant therewithin.

In some embodiments, the proximal end portion defines an opening for coupling a tilt control lead with the anchor. In some embodiments, the anchor is configured to be rotated when a force is exerted on the tilt control lead. In some embodiments, the anchor is rotated to align the first axis and the second axis for removal of the anchor and such that the distal end portion faces the passageway.

In some embodiments, the proximal end portion is substantially cylindrical. In some embodiments, the distal end portion is substantially conical. In some embodiments, the distal end portion defines an opening for engaging a delivery lead with the anchor. In some embodiments, the anchor is composed of at least one of a bio-compatible material, plastic, polypropylene, metal, ceramic, polymer, magnet, and alloy.

In some embodiments, the bodily implant is a mesh. In some embodiments, the bodily implant is composed of a bio-compatible material. In some embodiments, the bodily implant comprises at least one end portion, wherein the at least one end portion of the bodily implant is engaged with the anchor at the implant engaging portion.

In some embodiments, a method for anchoring a bodily implant within a body of a patient includes (1) inserting the bodily implant within the patient's body through a passageway, the passageway defining a first axis, an end portion of the bodily implant being coupled to a lateral portion of an anchor, the anchor defining a second axis extending from a distal end portion of the anchor to a proximal end portion of the anchor; and (2) exerting a force configured to rotate the anchor such that the second axis defined by the anchor forms an angle with the first axis defined by the passageway.

In some embodiments, the anchor is a first anchor and the end portion is a first end portion of the bodily implant. The method includes coupling a second end portion of the bodily implant to a second anchor at a lateral portion of the second anchor.

In some embodiments, the method includes inserting an elongated member into the patient's body to create the passageway therein. In some embodiments, the method includes exerting a force on a delivery lead coupled to the distal end portion of the anchor to align the second axis with the first axis. In some embodiments, the method includes exerting a force on a tilt control lead to rotate the anchor such that the second axis defined by the anchor is aligned with the first axis defined by the passageway.

While the invention has been disclosed in connection with the preferred embodiments shown and described in detail, various modifications and improvements thereon will become readily apparent to those skilled in the art. Accordingly, the spirit and scope of the present invention is not to be limited by the foregoing examples, but is to be understood in the broadest sense allowable by law.

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What is claimed is:

1. An anchor for anchoring a bodily implant within a body of a patient, the anchor comprising:
 - a distal end portion configured to pass through a passageway in the patient's body, the distal end portion having a lumen defined therethrough, the passageway defining a first axis;
 - a removable delivery lead extending through the lumen, the delivery lead configured to facilitate insertion of the anchor in the passageway;
 - a proximal end portion disposed longitudinally opposite to the distal end portion on the anchor, the proximal end portion including a tilt control lead,
 wherein the anchor defines a second axis extending from the distal end portion to the proximal end portion, the second axis being substantially parallel with the first axis defined by the passageway when the anchor is inserted in the passageway; and
 - a medial portion having an implant engaging portion for engaging the bodily implant, the implant engaging portion disposed on a lateral portion of the anchor, the implant engaging portion including a first protuberance and a second protuberance configured to engage the bodily implant, at least one of the first protuberance and the second protuberance being movable relative to a body of the anchor so as to engage the bodily implant, the tilt control lead extending from the proximal end portion, the tilt control lead being configured to facilitate rotation of the anchor in the passageway such that the second axis defined by the anchor is non-parallel with the first axis defined by the passageway to help secure the anchor in the passageway.
2. The anchor of claim 1, wherein at least one of the first protuberance and the second protuberance includes a movable locking mechanism for engaging the bodily implant.
3. The anchor of claim 1, wherein at least one of the first protuberance and the second protuberance includes a slidable locking mechanism for engaging the bodily implant.
4. The anchor of claim 1, wherein the first protuberance and the second protuberance are configured to interlock with each other.
5. The anchor of claim 1, wherein the proximal end portion defines an opening for coupling the tilt control lead with the anchor.
6. The anchor of claim 5, wherein the anchor is configured to be rotated when a force is exerted on the tilt control lead.
7. The anchor of claim 1, wherein the anchor is rotated to align the first axis and the second axis for removal of the anchor such that the distal end portion faces the passageway.
8. The anchor of claim 1, wherein the proximal end portion is substantially cylindrical.
9. The anchor of claim 1, wherein the distal end portion is substantially conical.
10. The anchor of claim 1, wherein the anchor is composed of at least one of a bio-compatible material, plastic, polypropylene, metal, ceramic, polymer, magnet, and alloy.
11. A medical device configured to be inserted within a body of a patient, the medical device comprising:
 - a bodily implant;
 - an anchor including:
 - a distal end portion configured to pass through a passageway in the patient's body, the passageway defining a first axis;
 - a proximal end portion disposed longitudinally opposite to the distal end portion of the anchor, the anchor defining a second axis extending from the distal end portion to the proximal end portion;

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- a medial portion having an implant engaging portion for engaging the bodily implant, the implant engaging portion disposed on a lateral portion of the anchor; and
- a lumen formed through the proximal portion; and
- a tilt control lead extending proximally from the lumen, wherein the tilt control lead, when manipulated by an operator, rotates the anchor from the proximal end portion such that the second axis defined by the anchor forms a non-zero angle with the first axis defined by the passageway to help secure the anchor in the passageway; and
- a delivery lead extending from the distal end portion, the delivery lead configured to be removeably coupled with the anchor and facilitate insertion of the anchor in the passageway.
- 12.** The medical device of claim **11**, wherein the implant engaging portion includes at least one protuberance for engaging the bodily implant therewith.
- 13.** The medical device of claim **11**, wherein the implant engaging portion includes a movable locking mechanism for engaging the bodily implant.
- 14.** The medical device of claim **11**, wherein the implant engaging portion includes a slidable locking mechanism for engaging the bodily implant.
- 15.** The medical device of claim **11**, wherein the implant engaging portion includes a first protuberance and a second protuberance configured to interlock with each other and engage the bodily implant therewithin.
- 16.** An anchor for anchoring a bodily implant within a body of a patient, the anchor comprising:
- a distal end portion configured to pass through a passageway in the patient's body, the passageway defining a first axis, the distal end portion having a lumen formed there-through;

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- a removable delivery lead extending through the lumen and away from the distal end portion, the delivery lead configured to facilitate insertion of the anchor in the passageway;
- a proximal end portion disposed longitudinally opposite to the distal end portion on the anchor,
- wherein the anchor defines a second axis extending from the distal end portion to the proximal end portion;
- a medial portion having an implant engaging portion for engaging the bodily implant, the implant engaging portion disposed on a lateral portion of the anchor and including at least one protuberance for engaging the bodily implant therewith, the at least one protuberance being aligned along the second axis, the implant engaging portion further including a movable locking mechanism for engaging the bodily implant; and
- a tilt control lead extending from the proximal end portion, the tilt control lead being configured to facilitate rotation of the anchor such that the second axis defined by the anchor forms a non-zero angle with the first axis defined by the passageway to help secure the anchor in the passageway.
- 17.** The anchor of claim **16**, wherein the at least one protuberance includes a first protuberance and a second protuberance configured to interlock with each other and engage the bodily implant.
- 18.** The anchor of claim **16**, wherein the proximal end portion defines an opening for coupling the tilt control lead with the anchor.
- 19.** The anchor of claim **16**, wherein the anchor is configured to be rotated when a force is exerted on the tilt control lead.

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